TOOTOPET TEDEOTIS

5

CLAIMS

What is claimed is:

/FL 4,2,1,33

- 1. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) contacting a 3-Isopropylmalate dehydratase polypeptide with said test compound; and
 - b) detecting the presence or absence of binding between a test compound and said 3-Isopropylmalate dehydratase polypeptide,

wherein binding indicates that said test compound is a candidate for an antibiotic.

- 2. The method of claim 1, wherein said 3-Isopropylmalate dehydratase polypeptide is a fungal 3-Isopropylmalate dehydratase polypeptide.
- 3. The method of claim-1, wherein said 3-Isopropylmalate dehydratase polypeptide is a *Magnaporthe* 3-Isopropylmalate dehydratase polypeptide.
- 4. The method of claim 1, wherein said 3-Isopropylmalate dehydratase polypeptide is SEQ ID NO: 3.
- 5. A method for determining whether the antibiotic candidate of claim 1 has antifungal activity, further comprising:
 - contacting a fungus or fungal cells with said antibiotic candidate and detecting the decrease in growth, viability, or pathogenicity of said fungus or fungal cells.

- 6. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) contacting a test compound with at least one polypeptide selected from the group consisting of: a polypeptide having at least ten consecutive amino acids of a fungal 3-Isopropylmalate dehydratase; a polypeptide having at least 50% sequence identity with a fungal 3-Isopropylmalate dehydratase; and a polypeptide having at least 10% of the activity of a fungal 3-Isopropylmalate dehydratase; and
 - b) detecting the presence and/or absence of binding between said test compound and said polypeptide,

wherein binding indicates that said test compound is a candidate for an antibiotic.

- 7. A method for determining whether the antibiotic candidate of claim 6 has antifungal activity, further comprising:
 - contacting a fungus or fungal cells with said antibiotic candidate and detecting a decrease in growth, viability, or pathogenicity of said fungus or fungal cells.
- 8. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) contacting 2-Isopropylmalate and H₂O with a 3-Isopropylmalate dehydratase;
 - b) contacting 2-Isopropylmalate and H₂O with 3-Isopropylmalate dehydratase and a test compound; and
- 20 c) determining the change in concentration for at least one of the following: 2-Isopropylmalate, H₂O, and/or 3-Isopropylmalate,
 - wherein a change in concentration for any of the above substances between steps (a) and (b) indicates that said test compound is a candidate for an antibiotic.

- 9. The method of claim 8, wherein said 3-Isopropylmalate dehydratase is a fungal 3-Isopropylmalate dehydratase.
- 5 10. The method of claim 8, wherein said 3-Isopropylmalate dehydratase is a Magnaporthe 3-Isopropylmalate dehydratase.
 - 11. The method of claim 8, wherein said 3-Isopropylmalate dehydratase is SEQ ID NO:

 3.
 - 12. A method for determining whether the antibiotic candidate of claim 8 has antifungal activity, further comprising:
 contacting a fungus or fungal cells with said antibiotic candidate and detecting a decrease in growth, viability, or pathogenicity of said fungus or fungal cells.
 - 13. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) contacting 3-Isopropylmalate with a 3-Isopropylmalate dehydratase;
 - b) contacting 3-Isopropylmalate with a 3-Isopropylmalate dehydratase and a test compound; and
 - c) determining the change in concentration for at least one of the following: 2-Isopropylmalate, H₂O, and/or 3-Isopropylmalate,

5

wherein a change in concentration for any of the above substances between steps (a) and (b) indicates that said test compound is a candidate for an antibiotic.

- 14. The method of claim 13, wherein said 3-Isopropylmalate dehydratase is a fungal 3-Isopropylmalate dehydratase.
- 15. The method of claim 13, wherein said 3-Isopropylmalate dehydratase is a *Magnaporthe* 3-Isopropylmalate dehydratase.
- 16. The method of claim 13, wherein said 3-Isopropylmalate dehydratase is SEQ ID NO:
 3.
- 17. A method for determining whether the antibiotic candidate of claim 13 has antifungal activity, further comprising:
- contacting a fungus or fungal cells with said antibiotic candidate and detecting a decrease in growth, viability, or pathogenicity of said fungus or fungal cells.
 - 18. A method for identifying a test compound as a candidate for an antibiotic, comprising:
- a) contacting 2-Isopropylmalate and H₂O with a polypeptide selected from the group consisting of: a polypeptide having at least 50% sequence identity with 3-Isopropylmalate dehydratase; a polypeptide having at least 50% sequence identity with a 3-Isopropylmalate dehydratase and having at least 10% of the activity

- thereof; and a polypeptide comprising at least 100 consecutive amino acids of a 3-Isopropylmalate dehydratase;
- b) contacting 2-Isopropylmalate and H₂O with said polypeptide and a test compound; and
- c) determining the change in concentration for at least one of the following: 2-Isopropylmalate, H₂O, and/or 3-Isopropylmalate,

wherein a change in concentration for any of the above substances between steps (a) and (b) indicates that said test compound is a candidate for an antibiotic.

- 19. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) contacting 3-Isopropylmalate with a polypeptide selected from the group consisting of: a polypeptide having at least 50% sequence identity with a 3-Isopropylmalate dehydratase; a polypeptide having at least 50% sequence identity with a 3-Isopropylmalate dehydratase and at least 10% of the activity thereof; and a polypeptide comprising at least 100 consecutive amino acids of a 3-Isopropylmalate dehydratase;
 - b) contacting 3-Isopropylmalate, with said polypeptide and a test compound; and
 - c) determining the change in concentration for at least one of the following: 2-Isopropylmalate, H₂O, and/or 3-Isopropylmalate,

wherein a change in concentration for any of the above substances between steps (a) and (b) indicates that said test compound is a candidate for an antibiotic.

- 20. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) measuring the expression of a 3-Isopropylmalate dehydratase in a cell, cells, tissue, or an organism in the absence of a test compound;
- b) contacting said cell, cells, tissue, or organism with said test compound and measuring the expression of said 3-Isopropylmalate dehydratase in said cell, cells, tissue, or organism; and
 - c) comparing the expression of 3-Isopropylmalate dehydratase in steps (a) and (b), wherein a lower expression in the presence of said test compound indicates that said test compound is a candidate for an antibiotic.
 - 21. The method of claim 20 wherein said cell, cells, tissue, or organism is, or is derived from a fungus.
- 15 22. The method of claim 20 wherein said cell, cells, tissue, or organism is, or is derived from a *Magnaporthe* fungus or fungal cell.
 - 23. The method of claim 20, wherein said 3-Isopropylmalate dehydratase is SEQ ID NO: 3.
 - 24. The method of claim-20, wherein the expression of 3-Isopropylmalate dehydratase is measured by detecting IPMD1 mRNA.

- 25. The method of claim 20, wherein the expression of 3-Isopropylmalate dehydratase is measured by detecting 3-Isopropylmalate dehydratase polypeptide.
- 26. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) providing cells having one form of a 3-Isopropylmalate dehydratase gene, and providing comparison cells having a different form of a 3-Isopropylmalate dehydratase gene; and
 - b) contacting said cells and said comparison cells with a test compound and determining the growth of said cells and comparison cells in the presence of the test compound,
 - wherein a difference in growth between said cells and said comparison cells in the presence of said compound indicates that said compound is a candidate for an antibiotic.
- 27. The method of claim 26 wherein the cells and the comparison cells are fungal cells.
- 28. The method of claim 26 wherein the cells and the comparison cells are *Magnaporthe* cells.
- 29. The method of claim 26 wherein said form and said comparison form of the 3 Isopropylmalate dehydratase are fungal 3-Isopropylmalate dehydratases.

- 30. The method of claim 26, wherein at least one of the forms is a *Magnaporthe* 3-Isopropylmalate dehydratase.
- 31. The method of claim 26 wherein said form and said comparison form of the 3-Isopropylmalate dehydratase are non-fungal 3-Isopropylmalate dehydratases.
- 32. The method of claim 26 wherein one form of the 3-Isopropylmalate dehydratase is a fungal 3-Isopropylmalate dehydratase, and the other form is a non-fungal 3-Isopropylmalate dehydratase.
- 33. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - a) providing cells having one form of a gene in the L-leucine biochemical and/or genetic pathway and providing comparison cells having a different form of said gene.
 - b) contacting said cells and said comparison cells with a said test compound,
 - c) determining the growth of said cells and said comparison cells in the presence of said test compound,
 - wherein a difference in growth between said cells and said comparison cells in the presence of said test compound indicates that said test compound is a candidate for an antibiotic.
- 15 34. The method of claim 33 wherein the cells and the comparison cells are fungal cells.

- 35. The method of claim 33 wherein the cells and the comparison cells are *Magnaporthe* cells.
- 36. The method of claim 33 wherein said form and said different form of the L-leucine biosynthesis gene are fungal L-leucine biosynthesis genes.
- 37. The method of claim 33, wherein at least one form is a *Magnaporthe* L-leucine biosynthesis gene.
- 38. The method of claim 33 wherein said form and said different form of the L-leucine biosynthesis genes are non-fungal L-leucine biosynthesis genes.
- 39. The method of claim 33 wherein one form of the L-leucine biosynthesis gene is a fungal L-leucine biosynthesis gene, and the different form is a non-fungal L-leucine biosynthesis gene.
- 40. A method for determining whether the antibiotic candidate of claim 33 has antifungal activity, further comprising:
- contacting a fungus or fungal cells with said antibiotic candidate and detecting a decrease in growth, viability, or pathogenicity of said fungus or fungal cells, wherein a decrease in growth, viability, or pathogenicity of said fungus or fungal cells indicates that the antibiotic candidate has antifungal activity.

5

- 41. A method for identifying a test compound as a candidate for an antibiotic, comprising:
 - (a) providing paired growth media; comprising a first medium and a second medium, wherein said second medium contains a higher level of L-leucine than said first medium;
 - (b) contacting an organism with a test compound;
 - (c) inoculating said first and said second media with said organism; and
 - (d) determining the growth of said organism,

wherein a difference in growth of the organism between said first and said second media indicates that said test compound is a candidate for an antibiotic.

- 42. The method of claim 41, wherein said organism is a fungus.
- 43. The method of claim 41, wherein said organism is Magnaporthe.
- 44. An isolated nucleic acid comprising a nucleotide sequence that encodes a polypeptide of SEQ ID NO: 3.
- 45. The nucleic acid of claim 44 comprising the nucleotide sequence of SEQ ID NO: 1.
- 46. An expression cassette comprising the nucleic acid of claim 45.
- 47. The isolated nucleic acid of claim 44 comprising a nucleotide sequence with at least 50 to at least 95% sequence identity to SEQ ID NO: 1.
- 48. A polypeptide consisting essentially of the amino acid sequence of SEQ ID NO: 3.
 - 49. A polypeptide comprising the amino acid sequence of SEQ ID NO: 3.